Amendments To The Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

In the Claims:

What is claimed is:

Claims 1-15. (Cancelled).

16. (New) A process for preparing (2S,3S,4R,5R)-2-(5-tert-butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol (Compound A) as Polymorph I comprising:

a) reacting 9-{(3aR,4R,6S,6aR)-6-[5-tert-butyl-1,3,4-oxadiazol-2-yl]-2,2-dimethyltetrahydrofuro[3,4-d][1,3]dioxol-4-yl}-N-(4-chloro-2-fluorophenyl)-9H-purin-6-amine with trifluoroacetic acid/water;

b) neutralizing with aqueous methanolic ammonia solution at 25-50 ^{o}C over at least one hour; and

c) cooling to 0-5 ^{o}C, optionally adding toluene.

17. (New) A process for preparing (2S,3S,4R,5R)-2-(5-tert-butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol (Compound A) as Polymorph I comprising:

a) dissolving Compound A in N,N-dimethylformamide and water wherein the N,N-dimethylformamide:water ratio is in the range 2.5:1 to 1.5:1 and the dilution is at least 15 volumes;

b) initiating crystallisation by either

adjusting the temperature to less than 25^{o}C; or

adjusting the temperature to less than 30^{o}C, and seeding with Polymorph I; and

c) optionally, adding toluene.

18. (New) A process for preparing (2S,3S,4R,5R)-2-(5-tert-butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol (Compound A) as Polymorph II comprising:

a) dissolving Compound A in N,N-dimethylformamide and water wherein the N,N-dimethylformamide:water ratio is in the range 2:1 to 1:2 and the dilution is at least 15 volumes;

b) initiating crystallisation by

adjusting the temperature to greater than 35°C; and optionally seeding with polymorph II; and

c) optionally, adding toluene.

19. (New) (2S,3S,4R,5R)-2-(5-tert-Butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol as Polymorph I in spheronised habit.

20. New) (2S,3S,4R,5R)-2-(5-tert-Butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol as Polymorph I in a habit obtainable by a process of claim 16.

21. (New) (2S,3S,4R,5R)-2-(5-tert-Butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol as Polymorph I in a habit obtainable by a process of claim 17.

22. (New) (2S,3S,4R,5R)-2-(5-tert-Butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol as Polymorph II in spheronised habit.

23. (New) (2S,3S,4R,5R)-2-(5-tert-Butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol as Polymorph II in a habit obtainable by a process of claim 18.

24. (New) A pharmaceutical composition comprising Polymorph I in a habit according to claim 19, and a pharmaceutically acceptable carrier and/or excipient.

25. (New) A pharmaceutical composition comprising Polymorph I in a habit according to claim 20, and a pharmaceutically acceptable carrier and/or excipient.

26. (New) A pharmaceutical composition comprising Polymorph I in a habit according to claim 21, and a pharmaceutically acceptable carrier and/or excipient.

27. (New) A pharmaceutical composition comprising Polymorph II in a habit according to claim 22, and a pharmaceutically acceptable carrier and/or excipient.

28. (New) A pharmaceutical composition comprising Polymorph II in a habit according to claim 23, and a pharmaceutically acceptable carrier and/or excipient.

29. (New) A method of treating a patient suffering from a condition where there is an advantage in decreasing plasma free fatty acid concentration, or reducing heart rate, or treating a patient suffering from ischemic heart disease, peripheral vascular disease, stroke, pain, CNS disorder, or sleep apnoea comprising administering a therapeutically effective amount of Polymorph I of (2S,3S,4R,5R)-2-(5-tert-butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol in a habit according to claim 19.

30. (New) A method of treating a patient suffering from a condition where there is an advantage in decreasing plasma free fatty acid concentration, or reducing heart rate, or treating a patient suffering from ischemic heart disease, peripheral vascular disease, stroke, pain, CNS disorder, or sleep apnoea

comprising administering a therapeutically effective amount of Polymorph I of (2S,3S,4R,5R)-2-(5-tert-butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol in a habit according to claim 20.

31. (New) A method of treating a patient suffering from a condition where there is an advantage in decreasing plasma free fatty acid concentration, or reducing heart rate, or treating a patient suffering from ischemic heart disease, peripheral vascular disease, stroke, pain, CNS disorder, or sleep apnoea comprising administering a therapeutically effective amount of Polymorph II of (2S,3S,4R,5R)-2-(5-tert-butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol in a habit according to claim 22.

32. (New) A method of treating a patient suffering from a condition where there is an advantage in decreasing plasma free fatty acid concentration, or reducing heart rate, or treating a patient suffering from ischemic heart disease, peripheral vascular disease, stroke, pain, CNS disorder, or sleep apnoea comprising administering a therapeutically effective amount of Polymorph II of (2S,3S,4R,5R)-2-(5-tert-butyl-[1,3,4]-oxadiazol-2-yl)-5-[6-(4-chloro-2-fluorophenylamino)-9H-purin-9-yl]-tetrahydrofuran-3,4-diol in a habit according to claim 23.